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Corrigendum

Corrigendum to "An Improved Stereocontrolled Synthesis of Pyochelin, Siderophore of *Pseudomonas aeruginosa* and *Burkholderia cepacia*" [Tetrahedron 56 (2000) 249]☆

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In the preparations of compounds **10** and **11**, the *t*-butyldiphenylsilyl group should be replaced by the *t*-butyldimethylsilyl group as follows:

Attempts to synthesise aldehyde 4 after protection of the phenol group as a *t*-butyldimethylsilyl ether

Methyl 2′-(2-*t*-butyldimethylsilyloxyphenyl)-2′-thia-zoline-4′-carboxylate 10. Imidazole (782 mg, 11.5 mmol) and *t*-butyldimethylsilyl chloride … 7.75 (dd; *J*=1.8, 7.8 Hz, 1H, H-6).

2′-(2-*t*-butyldimethylsilyloxyphenyl)-2′-thiazoline-4′-carboxaldehyde 11. To a solution of the former ester … 7.13–7.4 (m; 2H, H-4 & H-6).

The ^{1}H NMR and ^{13}C NMR spectra assignments of 2′-(2-hydroxyphenyl)-2′-thiazoline-4′-(*N*-methoxy, *N*-methyl) carboxamide **3c** should read as:

^{1}H NMR (200 MHz; $CDCl_3$): 3.30 (s; 3H, NCH_3); 3.48 (dd; *J*=9.2, 11 Hz, 1H, H-5′); 3.78 (t; *J*=9, 8 Hz; 1H, H-5′); 3.81 (s; 1H, OCH_3); 5.70 (t; *J*=8.8 Hz, 1H, H-4′); 6.88 (t; *J*=7.7 Hz, 1H, H-5); 6.98 (d; *J*=8.3 Hz, 1H, H-3); 7.34 (dd; *J*=1.4, 7.3 Hz, 1H, H-4); 7.42 (dt; *J*=1.6, 7.8 Hz, 1H, H-6).

^{13}C (100 MHz, $CDCl_3$): 32.6 (C-5′); 32.9 (NCH_3) 61.9 (OCH_3); 74.7 (C-4′); 116.2 (C-1); 117.2 (C-3); 119.0 (C-5); 130.6 (C-6); 133.5 (C-4); 159.2 (C-2), 169.2 (CO–N), 174.3 (C-2′).

The ^{1}H NMR and ^{13}C NMR spectra assignments of 2′-(2-hydroxyphenyl)-2′-thiazoline-4′-carboxylic acid **3a** should read as:

^{1}H NMR (200 MHz; $CDCl_3$): 3.69 (m; 2H, H-5′); 5.42 (t; *J*=8.12 Hz, 1H, H-4′); 6.89 (t; *J*=7.5 Hz, 1H, H-5); 7.03 (d; *J*=8.5 Hz, 1H, H-3); 7.43 (m; 2H, H-4 & H-6).

^{13}C NMR (100 MHz, $CDCl_3$): 33.6 (C-5′); 76.3 (C-4′); 115.8 (C-1); 117.4 (C-3); 119.1 (C-5); 130.8 (C-6); 133.8 (C-4); 159.1 (C-2), 174.7 (C-2′); 175.1 (COOH).

The ^{1}H NMR and ^{13}C NMR spectra assignments of methyl 2′-(2-hydroxyphenyl)-2′-thiazoline-4′-carboxylate **3b** should read as:

^{1}H NMR (200 MHz; $CDCl_3$): 3.58 (m; 2H, H-5′); 3.81 (s; 3H, OCH_3); 5.12 (t; *J*=9.28 Hz, 1H, H-4′); 6.87 (t; *J*=8.3 Hz, 1H, H-5); 6.97 (d; *J*=7.4 Hz, 1H, H-3); 7.28 (dd; *J*=1.3, 7.6 Hz, 1H, H-4); 7.78 (dt; *J*=1.3, 7.7 Hz, 1H, H-6).

^{13}C NMR (100 MHz, $CDCl_3$): 33.6 (C-5′); 52.8 (OCH_3); 76.6 (C-4′); 116.2 (C-1); 118.9(C-3); 119.1 (C-5); 130.6 (C-6); 133.5 (C-4); 159.2 (C-2); 174.3 (C-2′); 170.6 (COOMe).

The proton NMR data of aldehyde **4** should read as:

^{1}H NMR (200 MHz, $CDCl_3$): 3.39 (t; *J*=10 Hz, 1H, H-5′); 3.69 (dd, *J*=7.6, 11.4 Hz, 1H, H-5′); 5.33 (dd, *J*=7.8, 9.1 Hz, 1H, H-4′); 7.30–7.50 (m; 2H, H-4 & H-6), 6.80–7.05 (m; 2H, H-3 & H-5), 9.84 (s; 1H, aldehydic proton); 12.2 (s; 1H, O*H*).

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